Downloaded: 23/11/2024
Approved: 18/11/2024

Wiktoria Niewiadomska
Registration number: 220119496
Ophthalmology and Orthoptics
Programme: Vision and Strabismus MMedSci

Dear Wiktoria

PROJECT TITLE: Do Orthoptists advise near activities alongside amblyopia treatment? A survey of registered Orthoptists within the UK
APPLICATION: Reference Number 064957

On behalf of the University ethics reviewers who reviewed your project, I am pleased to inform you that on 18/11/2024 the above-named project was **approved** on ethics grounds, on the basis that you will adhere to the following documentation that you submitted for ethics review:

- University research ethics application form 064957 (form submission date: 02/11/2024); (expected project end date: 04/08/2025).
- Participant information sheet 1144815 version 2 (02/11/2024).
- Participant consent form 1144816 version 1 (21/10/2024).
- Participant consent form 1145212 version 1 (02/11/2024).

If during the course of the project you need to deviate significantly from the above-approved documentation please inform me since written approval will be required.

Your responsibilities in delivering this research project are set out at the end of this letter.

Yours sincerely

Kate Chadwick
Ethics Admin
School of Allied Health Professions, Nursing and Midwifery

Please note the following responsibilities of the researcher in delivering the research project:

- The project must abide by the University's Research Ethics Policy: https://www.sheffield.ac.uk/research-services/ethics-integrity/policy
- The project must abide by the University's Good Research & Innovation Practices Policy: https://www.sheffield.ac.uk/polopoly_fs/1.671066!/file/GRIPPolicy.pdf
- The researcher must inform their supervisor (in the case of a student) or Ethics Admin (in the case of a member of staff) of any significant changes to the project or the approved documentation.
- The researcher must comply with the requirements of the law and relevant guidelines relating to security and confidentiality of personal data.
- The researcher is responsible for effectively managing the data collected both during and after the end of the project in line with best practice, and any relevant legislative, regulatory or contractual requirements.